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II. Pending Claims

- 1. (Previously Presented) A pharmaceutical composition comprising:
- (a) a cyclodextrin; and
- (b) a lipidated glycopeptide antibiotic or a pharmaceutically acceptable salt thereof.
- (Previously Presented) The pharmaceutical composition of claim 1 which further comprises water.
 - 3. (Previously Presented) The pharmaceutical composition of claim 1 which is a powder.
- 4. (Previously Presented) The pharmaceutical composition of claim 1 which is a lyophilized powder.
 - 5. (Previously Presented) A pharmaceutical composition comprising:
 - (a) an aqueous cyclodextrin carrier and
- (b) a therapeutically effective amount of a lipidated glycopeptide antibiotic or a pharmaceutically acceptable salt thereof.
- 6. (Previously Presented) The pharmaceutical composition of Claim 5, wherein the pharmaceutical composition comprises:
 - a therapeutically effective amount of a lipidated glycopeptide antibiotic or a pharmaceutically acceptable salt thereof;
 - (b) 1 to 40 weight percent of a cyclodextrin; and
 - (c) 60 to 99 weight percent of water, provided that the components of the composition total 100 weight percent.
 - 7. (Original) The pharmaceutical composition of Claim 5, wherein the cyclodextrin is

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hydroxypropyl- β -cyclodextrin or sulfobutyl ether β -cyclodextrin.

- 8. (Original) The pharmaceutical composition of Claim 7, wherein the cyclodextrin is hydroxypropyl-β-cyclodextrin.
- 9. (Original) The pharmaceutical composition of Claim 6, wherein the cyclodextrin comprises about 5 to 35 weight percent of the composition.
- 10. (Original) The pharmaceutical composition of Claim 9, wherein the cyclodextrin comprises about 10 to 30 weight percent of the composition.
 - 11. (Canceled)
 - 12. (Canceled)
 - 13. (Canceled)
- 14. (Withdrawn) A method of treating a mammal having a bacterial disease, the method comprising administering to the mammal a pharmaceutical composition of claim 1.
- 15. (Withdrawn) A method of treating a bacterial disease in a mammal, the method comprising administering to the mammal a therapeutically effective amount of a lipidated glycopeptide antibiotic or a pharmaceutically acceptable salt thereof in combination with a cyclodextrin.

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- 16. (Withdrawn) A method for reducing tissue accumulation of a lipidated glycopeptide antibiotic when administered to a mammal, the method comprising administering the lipidated glycopeptide antibiotic to the mammal in a pharmaceutical composition comprising a cyclodextrin and a therapeutically effective amount of the lipidated glycopeptide antibiotic or a pharmaceutically acceptable salt thereof.
- 17. (Withdrawn) A method for reducing nephrotoxicity produced by a lipidated glycopeptide antibiotic when administered to a mammal, the method comprising administering the lipidated glycopeptide antibiotic to the mammal in a pharmaceutical composition comprising a cyclodextrin and a therapeutically effective amount of the lipidated glycopeptide antibiotic or a pharmaceutically acceptable salt thereof.
 - 18. (Canceled)
 - 19. (Canceled)